Amendments to the Claims:

- (currently amended) A method for enhancing preservative efficacy of [[a]] an ophthalmic composition, the composition comprising including a preservative effective amount and a soluble amount of a
- $\underline{0.001~wt\%~to~0.05~wt\%~of}~zinc, \ \mbox{eompound, wherein the composition has less than}$ a-preservative-effective amount of a primary

an organic, nitrogen-containing preservative agent,

a borate buffer, and

0 wt% to 0.01 wt% of a chelating agent, wherein the composition has an osmolality of from 225 mOsm.kg to 400 mOsm/kg.

Claims 2. - 5. Canceled

- (currently amended) The method of claim 5, wherein the polyeationic material is 1, further comprising a cationic cellulosic polymer.
- 7. (currently amended) The method of claim 6, wherein the composition has a minimum of about 0.001 wt.% and a maximum of about 0.5 wt.% cationic cellulosic polymer based upon the weight of the composition.

Claims 9. - 12. Canceled

13. (currently amended) The method of claim 1, further-comprising wherein the composition further comprises a therapeutically effective amount of therapeutic agent selected from the group comprising glaucoma agents, muscarinics, carbonic anhydrase inhibitors, dopaminergic agonists and antagonists, anti-infectives, non-steroidal and

steroidal anti-inflammatories, prostaglandins, enzymes, growth factors, anti-allergics, beta-blockers and mixtures and combinations thereof.

Claims 14. - 24. Canceled

- 25. (currently amended) The eomposition of claim 14, having method of claim 1, wherein the composition has the form of an eye drop solution.
- 26. (currently amended) The eomposition of claim 14, having method of claim 1, wherein the composition has the form of a contact lens treating solution.
- 27. (currently amended) The eomposition of claim 14, having method of claim 1, wherein the composition is suitable for direct instillation in the eye without causing irritation to eye tissue.

Claims 28. Canceled

29. (currently amended) An ophthalmic composition comprising: water:

at least one therapeutic agent in a therapeutically effective amount; and a preservative effective amount and a soluble amount of a zine compound 0.001 wt% to 0.05 wt% of zinc.

an organic, nitrogen-containing preservative agent,

a borate buffer, and

0 wt% to 0.01 wt% of a chelating agent, wherein the composition has an osmolality of from 225 mOsm/kg to 400 mOsm/kg.

Claims 30. - 34. Canceled

35. (currently amended) The composition of claim 34, wherein the cationic polysaceharide includes 29, further comprising a cationic cellulosic polymer.

- 36. (currently amended) The composition of claim [[34]] 35, wherein the composition has a minimum of about 0.001 wt.% and a maximum of about 0.5 wt.% of cationic cellulosic polymer based upon the total weight of the polymer.
- 37. (currently amended) The composition of claim [[33]] 36, wherein the polyeationic material cationic cellulosic polymer is Polymer JR.

Claims 38. - 39. Canceled

- 40. (original) The composition of claim 29, having the form of an eye drop solution.
- 41. (original) The composition of claim 29, having the form of a contact lens treating solution.
- 42. (original) The composition of claim 29, being suitable for direct instillation in the eye without irritation to eye tissue.
- 43. (currently amended) The composition of claim 29, wherein the <u>further comprising a</u> therapeutic agent is selected from the group consisting of glaucoma agents, muscarinics, carbonic anhydrase inhibitors, dopaminergic agonists and antagonists, anti-infectives, non-steroidal and steroidal anti-inflammatories, prostaglandins, enzymes, growth factors, anti-allergics, beta-blockers and mixtures and combinations thereof.
- 44. (currently amended) A method of treating an ophthalmic condition comprising administering a composition of claim 43 comprising water, a therapeutically effective amount of a therapeutic agent and a preservative-effective amount and a soluble amount of a zine compound.

Claims 45 - 78 Canceled